Adequate nutrition status important for bone mineral density improvement in a patient with anorexia nervosa

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Abstract: Low bone mineral density (BMD) is one of the most frequent complications of anorexia nervosa (AN). We report the clinical outcomes of a female patient with severe AN, whose chest had become deformed due to thoracic fracture. Lumbar BMD was 0.358 g/cm² (T-score = −6.3), and total hip BMD was 0.411 g/cm² (T-score = −4.4). Active vitamin D increased these parameters by 81.0% and 57.4%, respectively, but a drop in her nutrition status afterward resulted in a sharp decrease in BMD values. These findings suggest that adequate nutrient intake is essential for effective osteoporosis treatment in patients with AN.

Keywords: anorexia nervosa, bone mineral density, daily teriparatide, osteoporosis

Introduction
Anorexia nervosa (AN) is a severe eating disorder characterized by low body weight, intense fear of weight gain, and undue influence of weight and shape on self-evaluation.1 The overall age- and sex-adjusted incidence rate of AN is reportedly 4.1 per 100,000 person-years (95% CI: 2.4–5.9), with a female-to-male ratio of age-adjusted rates of 1.2:1.2 AN tends to manifest during adolescence3 and has the highest mortality rate of any psychiatric disorder, with no gold standard treatment4 and high therapy dropout and relapse rates.

One of the first treatment approaches that were based on Bruch’s observations was the “Maudsley model” of family therapy, in which the aim was to put patients in control of eating behavior.5 Approximately two-thirds of patients who receive this treatment as a form of early intervention show good recovery after 1 year, and this intervention is now recommended as a first-line treatment for adolescents with AN.6 However, it is important to note that this approach was not found to be effective if the illness had persisted for more than 3 years, or if the onset of the disorder occurred after the age of 18 years.

Numerous reports on AN and osteoporosis (OP) have surfaced over the past 3–4 decades. El Ghoch et al7 recently reviewed that diminished bone mineral density (BMD) was one of the most frequent medical complications in AN; nearly 85% of females with AN have very low BMD, and consequently, a sevenfold increase in the risk of spontaneous fracture compared with healthy controls. McAnarney et al8 described multiple rib fractures in a patient with AN as a result of forced vomiting in an individual with fragile bones. Khosla et al9 reported a preponderance of cancellous bone fractures in AN individuals (vertebral: 81%; rib: 37%; wrist: 13%), with 13% experiencing hip fracture.
The current first-line drugs for OP are bisphosphonates (BPs). Third-generation nitrogen-containing BPs inhibit farnesyl pyrophosphate synthetase in the mevalonate pathway in osteoclasts. Various other drugs have been described for patients with OP, such as the active vitamin D analog 1α (OH) vitamin D₃ (alfacalcidol; ALF), frequently used in Japan.

Concerning the treatment of adolescents with AN and OP, most methods tested (eg, hormone replacement, oral contraceptive pills, and BPs) have yielded only modest or negligible BMD improvements, with no data on the effectiveness of other strategies, such as physical activity intervention and/or the available nutritional supplementation (calcium, vitamin D, etc). The only promising pharmacological treatment to date has been physiological estrogen replacement by means of transdermal estradiol associated with cyclic progesterone, which, despite comparable weight gains, was associated with a significantly greater increase in spine and hip BMD than a placebo in non-severely underweight adolescents with AN. However, these findings require replication in patients with more severe malnutrition.

**Case report**

The patient was a 32-year-old female with a weight of 43 kg and a height of 157 cm. Since she has refused to know her body weight at all times, we have not measured it after she visited us for the first time. She had a medical history of pubic fracture and had been an outpatient at Shinshu University School of Medicine for 10 years following hospitalization for severe weight loss.

Her chief complaints were bilateral thoracic pain that had suddenly manifested 1 month prior and a common cold persisting for 2 months. The pain was obvious when taking a deep breath, coughing, or rolling over. The rib cage had become deformed to resemble the circumference of a barrel.

On presentation, her lumbar 1–4 BMD (L-BMD) was 0.358 g/cm² (T-score = −6.3), and bilateral total hip BMD (H-BMD) was 0.411 g/cm² (T-score = −4.4). Spinal plain radiographs showed no apparent fractures. Serum albumin was 4.3 g/dL, and 1,25(OH)₂D₃ was 35.1 pg/dL; both were within normal range. Serum bone alkaline phosphatase (BAP) was 212.0 U/L, and urinary N-terminal telopeptide of type I collagen was 226.0 nmol BCE/mmol Cr, which indicated extremely accelerated bone metabolism (Tables 1 and 2). She was diagnosed as having OP based on BMD measured by dual-energy X-ray absorption, for which ALF treatment was soon commenced. Thoracic pain subsided thereafter.

The values of bone turnover markers were gradually decreased after the therapy. At 3 years of treatment, alkaline phosphatase was 124 U/L (89.6% decrease from peak value), and BAP was 11.6 U/L (94.9% decrease from peak value). 25(OH)D and deoxypyridinoline values were slightly elevated, and tartrate-resistant acid phosphatase-5b was within normal range, showing no obvious acceleration of bone metabolism.

In 8 months of treatment, the percentage changes of L-BMD or H-BMD were increased to 70.7% or 41.4%, respectively. With respect to the peak values, L-BMD was 0.647 g/cm² (T-score = −2.4; 81.0% increase; Figure 1), and H-BMD was 0.648 g/cm² (T-score = −3.5; 57.4% increase; Figure 2).

She started to complain of deterioration of unbalanced diet and weight loss after the 2.5-year treatment. Her nutritional status soon degenerated, and serum albumin decreased to 3.4 g/dL. At 4 years of treatment, L-BMD had fallen to 0.484 g/cm² (T-score = −4.7; 25.2% decrease from peak value), and H-BMD was 0.605 g/cm² (T-score = −2.7; 6.6% decrease from peak value).

This patient gave written informed consent to publication of the patient’s personal medical information prior to her inclusion in this report.

**Discussion**

There have been numerous reports of accelerated bone resorption and inhibited bone formation in AN. In the current case, both bone resorption and bone formation were initially accelerated. We previously reported that bone turnover markers were significantly increased with accompanying back pain in elderly women, presumably due to insufficiency fracture, and that various bone fragility fractures might increase bone turnover markers. Our patient had suffered multiple fractures prior to her visit to our facility, and her thoracic circumference resembled that of a barrel. Thus, her fractures might have caused a remarkable increase in bone turnover markers. Despite the nonuse of bone anti-resorptive drugs, her BAP decreased from 228.0 to 11.6 U/L (94.9%) over 3 years, suggesting that bone metabolism improved from fracture healing. In patients with AN showing exceptional

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**Table 1 Changes in serum TP, Alb, and ALP**

<table>
<thead>
<tr>
<th>OP</th>
<th>0 M</th>
<th>2 M</th>
<th>1 Y</th>
<th>2 Y</th>
<th>2.5 Y</th>
<th>3 Y</th>
<th>Unit</th>
<th>Reference</th>
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</thead>
<tbody>
<tr>
<td>TP</td>
<td>6.6</td>
<td>6.5</td>
<td>6.5</td>
<td>6.8</td>
<td>5.9</td>
<td>4.9</td>
<td>g/dL</td>
<td>6.6–8.3</td>
</tr>
<tr>
<td>Alb</td>
<td>4.3</td>
<td>4.3</td>
<td>4.3</td>
<td>4.5</td>
<td>3.9</td>
<td>3.4</td>
<td>g/dL</td>
<td>3.7–4.9</td>
</tr>
<tr>
<td>ALP</td>
<td>1,187</td>
<td>1,092</td>
<td>307</td>
<td>215</td>
<td>177</td>
<td>124</td>
<td>U/L</td>
<td>105–368</td>
</tr>
</tbody>
</table>

**Abbreviations:** Alb, albumin; ALP, alkaline phosphatase; M, month(s); OP, observational period; TP, total protein; Y, year(s).
Adequate nutrition status important for BMI

Bone metabolism, especially the enhancement of bone formation marker, very severe AN complicated with fracture and pain is highly probable.

Evidence from a large number of studies has suggested that weight recovery in adolescents with AN may not be sufficient to fully reverse the detrimental effects of prolonged undernutrition on skeletal development. The benefits of exercise and calcium/vitamin D supplementation on BMD in patients with AN are also equivocal. Russell et al have reported that BPs can be an agent to treat AN except for premenopausal women and that the increase in L-BMD or H-BMD was 3%–4% or 2%, respectively. Isobe et al have recently reported that denosumab was effective for three patients with AN. In their report, the increases in L-BMD or H-BMD in those three cases were 15.7%, 18.6%, or none, and 35.2%, 11.6%, or 10.7%, respectively. Compared with those data, this study showed that the increase in L-BMD or H-BMD was 81% or 57.4% at the best during the study period, which showed a much better improvement than that in their study.

In this study, the increase in L-BMD was 70.7% and that in H-BMD was 41.4% at only 8 months of the treatment, which was too excellent considering her therapy being vitamin D alone. In our report on pregnancy and lactation-associated OP, we described two cases occurring in the early postpartum period that led to multiple spinal compression fractures. BMD gains were impressive in the early phases of treatment by combined vitamins D and K as follows: 19.7% at 1 year, 23.3% at 2 years, and 36.1% at 4 years in one case and 13.3% at 1 year, 17.3% at 2 years, and 26.3% at 3 years in the other (unpublished data). Thus, combination vitamin D and K therapy enabled a marked gradual increase in BMD in pregnancy and lactation-associated OP. In these cases, T-score of L-BMD was $-3.6$ SD and $-3.7$ SD, respectively, revealing that the BMD values were decreased markedly. Based on those findings and the results of the current case of AN, decreased BMD by repetitive fractures could be naturally recovered to some extent? We speculate that vitamin D and/or K may be instrumental in this process.

When our patient’s AN status deteriorated after year 3, serum albumin became decreased from 4.3 to 3.4 g/dL and

**Table 2** Changes in serum BAP, urinary NTX, I, serum 25(OH)D$_3$, 25(OH)D$_3$, DPD, ucOC, and TRACP-5b

<table>
<thead>
<tr>
<th></th>
<th>0 M</th>
<th>2 M</th>
<th>1 Y</th>
<th>2 Y</th>
<th>2.5 Y</th>
<th>3 Y</th>
<th>Unit</th>
<th>Reference</th>
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<tr>
<td>BAP</td>
<td>212.0</td>
<td>228.0</td>
<td>55.0</td>
<td>29.1</td>
<td>22.6</td>
<td>11.6</td>
<td>U/L</td>
<td>9.6–35.4</td>
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<td>Urinary NTX</td>
<td>226.0</td>
<td>276.2</td>
<td>152.9</td>
<td>92.5</td>
<td>118.2</td>
<td>–</td>
<td>nmol BCE/mmol Cr</td>
<td>95–54.3</td>
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<td>1,25(OH)$_2$D$_3$</td>
<td>–</td>
<td>35.1</td>
<td>43.3</td>
<td>–</td>
<td>–</td>
<td>–</td>
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<td>20–60</td>
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<tr>
<td>25(OH)D$_3$</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>5.0</td>
<td>–</td>
<td>ng/mL</td>
<td>7–41</td>
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<tr>
<td>DPD</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>11.7</td>
<td>–</td>
<td>nmol BCE/mmol Cr</td>
<td>2.8–7.6</td>
</tr>
<tr>
<td>ucOC</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>0.38</td>
<td>–</td>
<td>ng/mL</td>
<td>&lt;4.5</td>
</tr>
<tr>
<td>TRACP-5b</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>149</td>
<td>mU/dL</td>
<td>120–420</td>
</tr>
</tbody>
</table>

**Abbreviations:** BAP, bone alkaline phosphatase; DPD, deoxypyridinoline; M, month(s); NTX, N-terminal telopeptide of type-I collagen; TRACP, tartrate-resistant acid phosphatase; ucOC, undercarboxylated osteocalcin; Y, year(s).

![Figure 1](image1.png) **Figure 1** L-BMD (0.358 g/cm$^2$; T-score = −6.3) prior to treatment.

**Note:** At 3 years of therapy, it was 0.647 g/cm$^2$ (T-score = −2.4; 57.4% increase).

**Abbreviations:** BMD, bone mineral density; L-BMD, lumbar 1–4 BMD.

![Figure 2](image2.png) **Figure 2** Total hip BMD (0.411 g/cm$^2$; T-score = −4.4) prior to treatment.

**Note:** At 3 years of therapy, it was 0.648 g/cm$^2$ (T-score = −3.5; 81% increase).

**Abbreviation:** BMD, bone mineral density.
25(OH)D$_3$ remained at 5.0 ng/mL regardless of active vitamin D administration. Serum 25(OH)D$_3$ is the major and main storage form of vitamin D whose sufficiency level is estimated as 20–30 ng/mL. Therefore, 5.0 ng/dL represented a serious vitamin D deficiency, and BMD rapidly decreased irrespective of vitamin D continuation. Jáuregui-Lobera et al$^{17}$ reported that the most effective strategy to recover BMD in AN seems to be weight gain and menstrual recovery. Indeed, the improvement in nutrition may supersede all other treatment modalities for OP associated with AN.

Overall, it appeared that worsened AN causing inadequate nutrition and a rapid decline in BMD that was refractory to active vitamin D therapy may have increased our patient’s risk of fracture. Accordingly, anti-resorption drugs may be advocated for OP complicated with AN, especially when nutrition status may become compromised.

Disclosure

The authors report no conflicts of interest in this work.

References


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